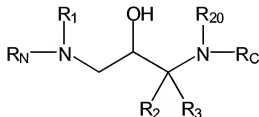


Listing of claims:

The following listing of claims will replace all prior versions and listings of claims in the application.

1. (Currently Amended) A compound of the formula I:



or a pharmaceutically acceptable salt or ester thereof, wherein R₂₀ is H, ~~C₁₋₆-alkyl or alkenyl, C₁₋₆-haloalkyl or C₄₋₆-cycloalkyl;~~ R₁ is ~~-(CH₂)₁₋₄-S(O)₀₋₂-(C₁₋₆-alkyl), or~~

~~C₁₋₁₀-alkyl optionally substituted with 1, 2, or 3 groups independently selected from halogen, -OH, -O-, -SH, -C≡N, -CF₃, -C₁₋₆-alkoxy, amino, mono- or dialkylamino, -N(R)C(O)R', -OC(=O)-amino and -OC(=O)-mono- or dialkylamino, or~~

~~C₂-C₆-alkenyl or C₂-C₆-alkynyl, each of which is optionally substituted with 1, 2, or 3 groups independently selected from halogen, -OH, -SH, -C≡N, -CF₃, C₁₋₆-alkoxy, amino, and mono- or dialkylamino, or~~

aryl, heteroaryl, heterocyclyl, -C₁₋₆ alkyl-aryl, -C₁₋₆ alkyl-heteroaryl, or -C₁₋₆ alkyl-heterocyclyl, where the ring portions of each are optionally substituted with 1, 2, 3, or 4 groups independently selected from halogen, -OH, -SH, -C≡N, -NR₁₀₅R'₁₀₅, -CO₂R, -N(R)COR', or -N(R)SO₂R', -C(=O)-(C₁₋₆) alkyl, -SO₂-amino, -SO₂-mono or dialkylamino, -C(=O)-amino, -C(=O)-mono or dialkylamino, -SO₂-(C₁₋₆) alkyl, or

C₁-C₆ alkoxy optionally substituted with 1, 2, or 3 groups which are independently selected from halogen, ~~or~~

~~C₃-C₆ cycloalkyl optionally substituted with 1, 2, or 3 groups independently selected from halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, amino, C₁-C₆ alkyl and mono- or dialkylamino, or~~

~~C₂-C₁₀ alkyl optionally substituted with 1, 2, or 3 groups independently selected from halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, amino, mono- or dialkylamino and C₁-C₆ alkyl, or~~

~~C₂-C₁₀ alkenyl or C₂-C₁₀ alkynyl each of which is optionally substituted with 1, 2, or 3 groups independently selected from halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, amino, C₁-C₆ alkyl and mono- or dialkylamino, and the heterocyclyl group is optionally further substituted with one;~~

R and R' independently are hydrogen, C₁-C₁₀ alkyl, C₁-C₁₀ alkylaryl or C₁-C₁₀ alkylheteroaryl;

R_C is hydrogen, -(CR₂₄₅R₂₅₀)₀₋₄-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-aryl-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-heterocyclyl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl-aryl, -[C(R₂₅₅)(R₂₆₀)]₁₋₃-CO-N-(R₂₅₅)₂, -CH(aryl)₂, -CH(heteroaryl)₂, -CH(heterocyclyl)₂, -CH(aryl)(heteroaryl), -(CH₂)₀₋₁-CH((CH₂)₀₋₆-OH)-(CH₂)₀₋₁-aryl, -(CH₂)₀₋₁-CH((CH₂)₀₋₆-OH)-(CH₂)₀₋₁-heteroaryl, -CH(-aryl or -heteroaryl)-CO-O(C₁-C₄ alkyl), -CH(-CH₂-OH)-CH(OH)-phenyl-NO₂, (C₁-C₆ alkyl)-O-(C₁-C₆ alkyl)-OH; -CH₂-NH-CH₂-CH(-O-CH₂-CH₃)₂, -(CH₂)₀₋₆-C(=NR₂₃₅)(NR₂₃₅R₂₄₀), or

~~C₁-C₁₀-alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, -OC-NR₂₃₅R₂₄₀, -S(=O)₀₋₂(C₁-C₆-alkyl), -SH, -NR₂₃₅-C-NR₂₃₅R₂₄₀, -C-NR₂₃₅R₂₄₀, and -S(=O)₀₋₂NR₂₃₅R₂₄₀, or -(CH₂)₀₋₅-(C₂-C₆) cycloalkyl wherein the cycloalkyl is optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, -CO₂H, and -CO₂-(C₁-C₄-alkyl), or~~

cyclopentyl, cyclohexyl, or cycloheptyl ring fused to aryl, heteroaryl, or heterocyclyl wherein one, two or three carbons of the cyclopentyl, cyclohexyl, or cycloheptyl is optionally replaced with a heteroatom independently selected from NH, NR₂₁₅, O, or S(=O)₀₋₂, and wherein the cyclopentyl, cyclohexyl, or cycloheptyl group can be optionally substituted with one or two groups that are independently R₂₀₅, =O, -CO-NR₂₃₅R₂₄₀, or -SO₂-(C₁-C₄ alkyl), or

~~C₂-C₁₀-alkenyl or C₂-C₁₀-alkynyl, each of which is optionally substituted with 1, 2, or 3 R₂₀₅-groups, wherein~~

each aryl and heteroaryl is optionally substituted with 1, 2, or 3 R₂₀₀, and wherein each heterocyclyl is optionally substituted with 1, 2, 3, or 4 R₂₁₀;

R₂₀₀ at each occurrence is independently selected from -OH, -NO₂, halogen, -CO₂H, C≡N, -(CH₂)₀₋₄-CO-NR₂₂₀R₂₂₅, -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl), -(CH₂)₀₋₄-CO-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-CO-aryl, -(CH₂)₀₋₄-CO-heteroaryl, -(CH₂)₀₋₄-CO-heterocyclyl, -(CH₂)₀₋₄-CO-O-R₂₁₅, -(CH₂)₀₋₄-SO₂-NR₂₂₀R₂₂₅, -(CH₂)₀₋₄-SO-(C₁-C₈ alkyl), -(CH₂)₀₋₄-SO₂-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-SO₂-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-N(H or R₂₁₅)-CO-O-R₂₁₅, -(CH₂)₀₋₄-N(H or R₂₁₅)-CO-N(R₂₁₅)₂, -(CH₂)₀₋₄-N-CS-N(R₂₁₅)₂, -(CH₂)₀₋₄-N(H or R₂₁₅)-CO-R₂₂₀, -(CH₂)₀₋₄-NR₂₂₀R₂₂₅, -(CH₂)₀₋₄-O-CO-(C₁-C₆ alkyl), -(CH₂)₀₋₄-O-P(O)-(OR₂₄₀)₂, -(CH₂)₀₋₄-O-CO-N(R₂₁₅)₂, -(CH₂)₀₋₄-O-CS-N(R₂₁₅)₂,

$-(CH_2)_{0-4}-O-(R_{215})$, $-(CH_2)_{0-4}-O-(R_{215})-COOH$, $-(CH_2)_{0-4}-S-(R_{215})$,
 $-(CH_2)_{0-4}-O-(C_1-C_6 \text{ alkyl optionally substituted with 1, 2, 3, or 5 -F})$, $C_3-C_7 \text{ cycloalkyl}$, $-(CH_2)_{0-4}-N(H \text{ or } R_{215})-SO_2-R_{220}$,
 $-(CH_2)_{0-4}-C_3-C_7 \text{ cycloalkyl}$, or
 $C_1-C_{10} \text{ alkyl optionally substituted with 1, 2, or 3 } R_{205} \text{ groups}$, or
 $C_2-C_{10} \text{ alkenyl or } C_2-C_{10} \text{ alkynyl}$, each of which is optionally substituted with 1 or 2 R_{205} groups, wherein
the aryl and heteroaryl groups at each occurrence are optionally substituted with 1, 2, or 3 groups that are independently R_{205} , R_{210} , or
 $C_1-C_6 \text{ alkyl substituted with 1, 2, or 3 groups that are independently } R_{205} \text{ or } R_{210}$, and wherein
the heterocyclyl group at each occurrence is optionally substituted with 1, 2, or 3 groups that are independently R_{210} ;
 R_{205} at each occurrence is independently selected from $C_1-C_6 \text{ alkyl}$, halogen, $-OH$, $-O\text{-phenyl}$, $-SH$, $-C\equiv N$, $-CF_3$, $C_1-C_6 \text{ alkoxy}$, NH_2 , $NH(C_1-C_6 \text{ alkyl})$ or $N-(C_1-C_6 \text{ alkyl})(C_1-C_6 \text{ alkyl})$;
 R_{210} at each occurrence is independently selected from halogen, $C_1-C_6 \text{ alkoxy}$, $C_1-C_6 \text{ haloalkoxy}$, $-NR_{220}R_{225}$, OH , $C\equiv N$, $-CO-(C_1-C_4 \text{ alkyl})$, $-SO_2-NR_{235}R_{240}$, $-CO-NR_{235}R_{240}$, $-SO_2-(C_1-C_4 \text{ alkyl})$, $=O$, or $C_1-C_6 \text{ alkyl}$, $C_2-C_6 \text{ alkenyl}$, $C_2-C_6 \text{ alkynyl}$ or $C_3-C_7 \text{ cycloalkyl}$, each of which is optionally substituted with 1, 2, or 3 R_{205} groups;
 R_{215} at each occurrence is independently selected from $C_1-C_6 \text{ alkyl}$, $-(CH_2)_{0-2}-(\text{aryl})$, $C_2-C_6 \text{ alkenyl}$, $C_2-C_6 \text{ alkynyl}$, $C_3-C_7 \text{ cycloalkyl}$, and $-(CH_2)_{0-2}-(\text{heteroaryl})$, $-(CH_2)_{0-2}-(\text{heterocyclyl})$, wherein
the aryl group at each occurrence is optionally substituted with 1, 2, or 3 groups that are independently R_{205} or R_{210} , and wherein

the heterocyclyl and heteroaryl groups at each occurrence are optionally substituted with 1, 2, or 3 R₂₁₀;

R₂₂₀ and R₂₂₅ at each occurrence are independently selected from -H, -C₃-C₇ cycloalkyl, -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl), -(C₁-C₆ alkyl)-O-(C₁-C₃ alkyl), -C₂-C₆ alkenyl, -C₂-C₆ alkynyl, -C₁-C₆ alkyl chain with one double bond and one triple bond, -aryl, -heteroaryl, and -heterocyclyl, or -C₁-C₁₀ alkyl optionally substituted with -OH, -NH₂ or halogen, wherein

the aryl, heterocyclyl and heteroaryl groups at each occurrence are optionally substituted with 1, 2, or 3 R₂₇₀ groups

R₂₃₅ and R₂₄₀ at each occurrence are independently H, or C₁-C₆ alkyl;

R₂₄₅ and R₂₅₀ at each occurrence are independently selected from -H, C₁-C₄ alkyl, C₁-C₄ alkylaryl, C₁-C₄ alkylheteroaryl, C₁-C₄ hydroxyalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, -(CH₂)₀₋₄-C₃-C₇ cycloalkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, and phenyl; or

R₂₄₅ and R₂₅₀ are taken together with the carbon to which they are attached to form a carbocycle of 3, 4, 5, 6, or 7 carbon atoms, where one carbon atom is optionally replaced by a heteroatom selected from -O-, -S-, -SO₂-, and -NR₂₂₀-;

R₂₅₅ and R₂₆₀ at each occurrence are independently selected from -H, -(CH₂)₁₋₂-S(O)₀₋₂-(C₁-C₆ alkyl), -(C₁-C₄ alkyl)-aryl, -(C₁-C₄ alkyl)-heteroaryl, -(C₁-C₄ alkyl)-heterocyclyl, -aryl, -heteroaryl, -heterocyclyl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-aryl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-heteroaryl, -(CH₂)₁₋₄-R₂₆₅-(CH₂)₀₋₄-heterocyclyl, or

C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl or -(CH₂)₀₋₄-C₃-C₇ cycloalkyl, each of which is optionally substituted with 1, 2, or 3 R₂₀₅ groups, wherein

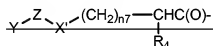
each aryl or phenyl is optionally substituted with 1, 2, or 3 groups that are independently R₂₀₅, R₂₁₀, or

C₁-C₆ alkyl substituted with 1, 2, or 3 groups that are independently R₂₀₅ or R₂₁₀, and wherein each heterocyclyl is optionally substituted with 1, 2, 3, or 4 R₂₁₀;

R₂₆₅ at each occurrence is independently -O-, -S- or -N(C₁-C₆ alkyl)-;

R₂₇₀ at each occurrence is independently R₂₀₅, halogen C₁-C₆ alkoxy, C₁-C₆ haloalkoxy, NR₂₃₅R₂₄₀, -OH, -C≡N, -CO-(C₁-C₄ alkyl), -SO₂-NR₂₃₅R₂₄₀, -CO-NR₂₃₅R₂₄₀, -SO₂-(C₁-C₄ alkyl), =O, or C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl or -(CH₂)₀₋₄-C₃-C₇ cycloalkyl, each of which is optionally substituted with 1, 2, or 3 R₂₀₅ groups;

R_N is R'₁₀₀, -C(=O)-NR'₁₀₀-R'₁₀₀, -C(=O)O-R'₁₀₀, -SO₂R'₁₀₀, -(CRR')₁₋₆R'₁₀₀, -C(=O)-(CRR')₀₋₆R₁₀₀, -C(=O)-(CRR')₁₋₆O-R'₁₀₀, -C(=O)-(CRR')₁₋₆-S-R'₁₀₀, -C(=O)-(CRR')₁₋₆-C(=O)-R₁₀₀, -C(=O)-(CRR')₁₋₆-SO₂-R₁₀₀, or -C(=O)-(CRR')₁₋₆-NR₁₀₀-R'₁₀₀;~~or~~



wherein

~~R₄ is selected from the group consisting of H; NH₂; -NH-(CH₂)_{n6}-R₄₋₁; -NHR₄₋₁; -NR₄₋₆C(O)R₄₋₁; C₁-C₄ alkyl-NHC(O)R₄₋₁; -(CH₂)₀₋₄R₄₋₁; -O-C₁-C₄-alkanoyl; OH; C₆-C₁₀-aryloxy optionally substituted with 1, 2, or 3 groups that are independently halogen, C₁-C₄ alkyl, -CO₂H, -C(O)-C₁-C₄-alkoxy, or C₁-C₄-alkoxy; C₁-C₆ alkoxy; aryl C₁-C₄-alkoxy; -NR₄₋₆CO₂R₄₋₁; -C₁-C₄ alkyl-NR₄₋₆CO₂R₄₋₁; -C≡N; -CF₂; -CF₂-CF₂; -C≡CH; -CH₂-CH=CH₂; -(CH₂)₁₋₄-R₄₋₁; -(CH₂)₁₋₄-NH-R₄₋₁; -O-(CH₂)_{n6}-R₄₋₁; -S-(CH₂)_{n6}-R₄₋₁; -(CH₂)₀₋₄-NHC(O)-(CH₂)₀₋₆-R₄₋₂; -(CH₂)₀₋₄-R₄₋₂-(CH₂)₀₋₄-R₄₋₁~~

wherein

~~n₆ is 0, 1, 2, or 3;~~

~~n₇ is 0, 1, 2, or 3;~~

~~R₄₋₁ is selected from the group consisting of SO₂-(C₁-C₆ alkyl), SO-(C₁-C₆ alkyl), S-(C₁-C₆ alkyl), S-CO-(C₁-C₆ alkyl), SO₂-NR₄₋₂R₄₋₃, CO-C₁-C₂ alkyl, CO-NR₄₋₃R₄₋₄, R₄₋₂ and R₄₋₃ are independently H, C₁-C₃ alkyl, or C₅-C₆ cycloalkyl, R₄₋₄ is alkyl, arylalkyl, alkanoyl, or arylalkanoyl, R₄₋₅ is H or C₁-C₆ alkyl,~~

~~— R₅ is selected from the group consisting of C₂-C₄ cycloalkyl, C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups that are independently halogen, NR₆R₇, C₁-C₄ alkoxy, C₅-C₆ heterocycloalkyl, C₅-C₆ heteroaryl, C₆-C₁₀ aryl, C₅-C₆ cycloalkyl C₁-C₄ alkyl, S-C₁-C₄ alkyl, SO₂-C₁-C₄ alkyl, CO₂H, CONR₆R₇, CO₂-C₁-C₄ alkyl, C₆-C₁₀ aryloxy, heteroaryl optionally substituted with 1, 2, or 3 groups that are independently C₁-C₄ alkyl, C₁-C₄ alkoxy, halogen, C₁-C₄ haloalkyl, or OH, heterocycloalkyl optionally substituted with 1, 2, or 3 groups that are independently C₁-C₄ alkyl, C₁-C₄ alkoxy, halogen, or C₂-C₄ alkanoyl, aryl optionally substituted with 1, 2, 3, or 4 groups that are independently halogen, OH, C₁-C₄ alkyl, C₁-C₄ alkoxy, or C₁-C₄ haloalkyl, and NR₆R₇, wherein~~

~~— R₆ and R₇ are independently selected from the group consisting of H, C₁-C₆ alkyl, C₅-C₆ alkanoyl, phenyl, SO₂-C₁-C₄ alkyl, phenyl C₁-C₄ alkyl,~~

~~R₈ is selected from the group consisting of SO₂-heteroaryl, SO₂-aryl, SO₂-heterocycloalkyl, SO₂-C₁-C₁₀ alkyl, C(O)NHR₉, heterocycloalkyl, S-C₁-C₆ alkyl, S-C₂-C₄ alkanoyl, wherein~~

~~— R₉ is aryl C₁-C₄ alkyl, C₁-C₆ alkyl, or H,~~

~~R₁₀ is H or C₁-C₆ alkyl,~~

~~R₁₁ is selected from the group consisting of aryl C₁-C₄ alkyl, C₁-C₆ alkyl optionally substituted with 1, 2, or~~

3 groups that are independently halogen, cyano, heteroaryl, NR_aR_b , $\text{C}(\text{O})\text{NR}_a\text{R}_b$, $\text{C}_3\text{-C}_4$ cycloalkyl, or $\text{C}_1\text{-C}_4$ alkoxy; heterocycloalkyl optionally substituted with 1 or 2 groups that are independently $\text{C}_1\text{-C}_4$ alkyl, $\text{C}_1\text{-C}_4$ alkoxy, halogen, $\text{C}_3\text{-C}_4$ alkanoyl, aryl $\text{C}_1\text{-C}_4$ alkyl, and $\text{SO}_2\text{-C}_1\text{-C}_4$ alkyl; alkenyl; alkynyl; heteroaryl optionally substituted with 1, 2, or 3 groups that are independently OH, $\text{C}_1\text{-C}_4$ alkyl, $\text{C}_1\text{-C}_4$ alkoxy, halogen, NH_2 , $\text{NH}(\text{C}_1\text{-C}_4\text{ alkyl})$ or $\text{N}(\text{C}_1\text{-C}_4\text{ alkyl})(\text{C}_1\text{-C}_4\text{ alkyl})$; heteroarylalkyl optionally substituted with 1, 2, or 3 groups that are independently $\text{C}_1\text{-C}_4$ alkyl, $\text{C}_1\text{-C}_4$ alkoxy, halogen, NH_2 , $\text{NH}(\text{C}_1\text{-C}_4\text{ alkyl})$ or $\text{N}(\text{C}_1\text{-C}_4\text{ alkyl})(\text{C}_1\text{-C}_4\text{ alkyl})$; aryl, heterocycloalkyl, $\text{C}_3\text{-C}_4$ cycloalkyl, and cycloalkylalkyl; wherein the aryl, heterocycloalkyl, $\text{C}_3\text{-C}_4$ cycloalkyl, and cycloalkylalkyl groups are optionally substituted with 1, 2, 3, 4 or 5 groups that are independently halogen, CN, NO_2 , $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy, $\text{C}_2\text{-C}_6$ alkanoyl, $\text{C}_1\text{-C}_6$ haloalkyl, $\text{C}_1\text{-C}_6$ haloalkoxy, hydroxy, $\text{C}_1\text{-C}_6$ hydroxyalkyl, $\text{C}_1\text{-C}_6$ alkoxy $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ thioalkoxy, $\text{C}_1\text{-C}_6$ thioalkoxy $\text{C}_1\text{-C}_6$ alkyl, or $\text{C}_1\text{-C}_6$ alkoxy $\text{C}_3\text{-C}_6$ alkoxy;

$\text{R}_{a,b}$ is heterocycloalkyl, heteroaryl, aryl, cycloalkyl, $\text{S}(\text{O})_{0-2}\text{-C}_1\text{-C}_6$ alkyl, CO_2H , $\text{C}(\text{O})\text{NH}_2$, $\text{C}(\text{O})\text{NH}(\text{alkyl})$, $\text{C}(\text{O})\text{N}(\text{alkyl})(\text{alkyl})$, $\text{CO}_2\text{-alkyl}$, $\text{NHS}(\text{O})_{0-2}\text{-C}_1\text{-C}_6$ alkyl, $\text{N}(\text{alkyl})\text{S}(\text{O})_{0-2}\text{-C}_1\text{-C}_6$ alkyl, $\text{S}(\text{O})_{0-2}$ heteroaryl, $\text{S}(\text{O})_{0-2}$ aryl, $\text{NH}(\text{arylalkyl})$, $\text{N}(\text{alkyl})(\text{arylalkyl})$, thioalkoxy, or alkoxy, each of which is optionally substituted with 1, 2, 3, 4, or 5 groups that are independently alkyl, alkoxy, thioalkoxy, halogen, haloalkyl, haloalkoxy, alkanoyl, NO_2 , CN, alkoxy carbonyl, or aminocarbonyl;

~~R_{3a} is absent, -O-, -C(O)-, -NH-, -N(alkyl)-, -NH-S(O)₀₋₂-,
-N(alkyl)-S(O)₀₋₂-, -S(O)₀₋₂-NH-, -S(O)₀₋₂-N(alkyl)-,
-NH-C(S)-, or -N(alkyl)-C(S)-;~~

~~R_{3b} is heteroaryl, aryl, arylalkyl, heterocycloalkyl, CO₂H,
-CO₂-alkyl, -C(O)NH(alkyl), -C(O)N(alkyl)-(alkyl),
-C(O)NH₂, C₁-C₆-alkyl, OH, aryloxy, alkoxy, arylalkoxy,
NH₂, NH(alkyl), N(alkyl)-(alkyl), or -C₁-C₆-alkyl-CO₂-
C₁-C₆-alkyl, each of which is optionally substituted
with 1, 2, 3, 4, or 5 groups that are independently
alkyl, alkoxy, CO₂H, -CO₂-alkyl, thioalkoxy, halogen,
haloalkyl, haloalkoxy, hydroxyalkyl, alkanoyl, NO₂, CN,
alkoxycarbonyl, or aminocarbonyl;~~

~~X' is selected from the group consisting of -C₁-C₆-alkylidenyl
optionally optionally substituted with 1, 2, or 3 methyl
groups, and -NR₄₋₆-; or~~

~~R₄ and R₄₋₆ combine to form -(CH₂)_{n±0}-, wherein
n±0 is 1, 2, 3, or 4;~~

~~Z is selected from the group consisting of a bond, SO₂, SO, S,
and C(O);~~

~~Y is selected from the group consisting of H, C₁-C₄-haloalkyl, C₂-
C₆-heterocycloalkyl, C₆-C₁₀-aryl, OH, -N(Y_a)(Y_b), C₁-C₁₀-alkyl
optionally substituted with 1 thru 3 substituents which can
be the same or different and are selected from the group
consisting of halogen, hydroxy, alkoxy, thioalkoxy, and
haloalkoxy, C₂-C₆-cycloalkyl optionally substituted with 1,
2, or 3 groups independently selected from C₁-C₂-alkyl, and
halogen, alkoxy, aryl optionally substituted with halogen,
alkyl, alkoxy, CN or NO₂, arylalkyl optionally substituted
with halogen, alkyl, alkoxy, CN or NO₂, wherein~~

~~Y_a and Y_b are the same or different and are H, C₁-C₁₀-alkyl
optionally substituted with 1, 2, or 3 substituents
selected from the group consisting of halogen, C₁-C₄-
alkoxy, C₂-C₆-cycloalkyl, and OH, C₂-C₆-alkenyl, C₂-C₆-~~

~~alkanoyl, phenyl, -SO₂-C₁-C₄-alkyl, phenyl-C₁-C₄-alkyl, or C₁-C₄-cycloalkyl-C₁-C₄-alkyl, or~~
~~Y₁, Y₂ and the nitrogen to which they are attached form a ring selected from the group consisting of piperazinyl, piperidinyl, morpholinyl, and pyrrolidinyl, wherein each ring is optionally substituted with 1, 2, 3, or 4 groups that are independently C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkoxy-C₁-C₄-alkyl, or halogen,~~
R₁₀₀ and R'₁₀₀ independently represent aryl, heteroaryl, -aryl-W-aryl, -aryl-W-heteroaryl, -aryl-W-heterocyclyl, -heteroaryl-W-aryl, -heteroaryl-W-heteroaryl, -heteroaryl-W-heterocyclyl, -heterocyclyl-W-aryl, -heterocyclyl-W-heteroaryl, -heterocyclyl-W-heterocyclyl, -CH[(CH₂)₀₋₂-O-R₁₅₀]- (CH₂)₀₋₂-aryl, -CH[(CH₂)₀₋₂-O-R₁₅₀]- (CH₂)₀₋₂-heterocyclyl or -CH[(CH₂)₀₋₂-O-R₁₅₀]- (CH₂)₀₋₂-heteroaryl, where the ring portions of each are optionally substituted with 1, 2, or 3 groups independently selected from
-OR, -NO₂, halogen, -C≡N, -OCF₃, -CF₃, -(CH₂)₀₋₄-O-P(=O)(OR)(OR'), -(CH₂)₀₋₄-CO-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-O-(CH₂)₀₋₄-CONR₁₀₂R'₁₀₂', -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl), -(CH₂)₀₋₄-CO-(CH₂)₀₋₄-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-R₁₁₀, -(CH₂)₀₋₄-R₁₂₀, -(CH₂)₀₋₄-R₁₃₀, -(CH₂)₀₋₄-CO-R₁₁₀, -(CH₂)₀₋₄-CO-R₁₂₀, -(CH₂)₀₋₄-CO-R₁₃₀, -(CH₂)₀₋₄-CO-R₁₄₀, -(CH₂)₀₋₄-CO-O-R₁₅₀, -(CH₂)₀₋₄-SO₂-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-SO-(C₁-C₈ alkyl), -(CH₂)₀₋₄-SO₂-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-SO₂-(CH₂)₀₋₄-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-N(R₁₅₀)-CO-O-R₁₅₀, -(CH₂)₀₋₄-N(R₁₅₀)-CO-N(R₁₅₀)₂, -(CH₂)₀₋₄-N(R₁₅₀)-CS-N(R₁₅₀)₂, -(CH₂)₀₋₄-N(R₁₅₀)-CO-R₁₀₅, -(CH₂)₀₋₄-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-R₁₄₀, -(CH₂)₀₋₄-O-CO-(C₁-C₆ alkyl), -(CH₂)₀₋₄-O-P(O)-(O-R₁₁₀)₂, -(CH₂)₀₋₄-O-CO-N(R₁₅₀)₂, -(CH₂)₀₋₄-O-CS-N(R₁₅₀)₂, -(CH₂)₀₋₄-O-(R₁₅₀)₂, -(CH₂)₀₋₄-O-R₁₅₀', -COOH, -(CH₂)₀₋₄-S-(R₁₅₀)₂, -(CH₂)₀₋₄-N(R₁₅₀)-SO₂-R₁₀₅,

$-(CH_2)_{0-4}-$ C_3-C_7 cycloalkyl, (C_2-C_{10}) alkenyl, or (C_2-C_{10}) alkynyl, or
 R_{100} is C_1-C_{10} alkyl optionally substituted with 1, 2, or 3 R_{115} groups, or
 R_{100} is $-(C_1-C_6 \text{ alkyl})-O-C_1-C_6 \text{ alkyl}$ or $-(C_1-C_6 \text{ alkyl})-S-(C_1-C_6 \text{ alkyl})$, each of which is optionally substituted with 1, 2, or 3 R_{115} groups, or
 R_{100} is C_3-C_8 cycloalkyl optionally substituted with 1, 2, or 3 R_{115} groups;
 W is $-(CH_2)_{0-4}-$, $-O-$, $-S(O)_{0-2}-$, $-N(R_{135})-$, $-CR(OH)-$ or $-C(O)-$;
 R_{102} and R_{102}' independently are hydrogen, or
 C_1-C_{10} alkyl optionally substituted with 1, 2, or 3 groups that are independently halogen, aryl or $-R_{110}$;
 R_{105} and R'_{105} independently represent $-H$, $-R_{110}$, $-R_{120}$, C_3-C_7 cycloalkyl, $-(C_1-C_2 \text{ alkyl})-(C_3-C_7 \text{ cycloalkyl})$, $-(C_1-C_6 \text{ alkyl})-O-(C_1-C_3 \text{ alkyl})$, C_2-C_6 alkenyl, C_2-C_6 alkynyl, or C_1-C_6 alkyl chain with one double bond and one triple bond, or
 C_1-C_6 alkyl optionally substituted with $-OH$ or $-NH_2$; or,
 C_1-C_6 alkyl optionally substituted with 1, 2, or 3 groups independently selected from halogen, or
 R_{105} and R'_{105} together with the atom to which they are attached form a 3 to 7 membered carbocyclic ring, where one member is optionally a heteroatom selected from $-O-$, $-S(O)_{0-2}-$, $-N(R_{135})-$, the ring being optionally substituted with 1, 2 or three R_{140} groups;
 R_{115} at each occurrence is independently halogen, $-OH$, $-CO_2R_{102}$, $-C_1-C_6$ thioalkoxy, $-CO_2$ -phenyl, $-NR_{105}R'_{135}$, $-SO_2-(C_1-C_8 \text{ alkyl})$, $-C(=O)R_{180}$, R_{180} , $-CONR_{105}R'_{105}$, $-SO_2NR_{105}R'_{105}$, $-NH-CO-(C_1-C_6 \text{ alkyl})$, $-NH-C(=O)-OH$, $-NH-C(=O)-OR$, $-NH-C(=O)-O$ -phenyl, $-O-C(=O)-(C_1-C_6 \text{ alkyl})$, $-O-C(=O)$ -amino, $-O-C(=O)$ -mono- or dialkylamino, $-O-C(=O)$ -phenyl, $-O-(C_1-C_6 \text{ alkyl})-CO_2H$, $-NH-SO_2-(C_1-C_6 \text{ alkyl})$, C_1-C_6 alkoxy or C_1-C_6 haloalkoxy;

- R₁₃₅ is C₁-C₆ alkyl, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₃-C₇ cycloalkyl, - (CH₂)₀₋₂- (aryl), - (CH₂)₀₋₂- (heteroaryl), or - (CH₂)₀₋₂- (heterocyclyl);
- R₁₄₀ is heterocyclyl optionally substituted with 1, 2, 3, or 4 groups independently selected from C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, hydroxy, cyano, nitro, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ haloalkyl, C₁-C₆ haloalkoxy, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino(C₁-C₆)alkyl, di(C₁-C₆)alkylamino(C₁-C₆)alkyl, and =O;
- R₁₅₀ is hydrogen, C₃-C₇ cycloalkyl, -(C₁-C₂ alkyl)-(C₃-C₇ cycloalkyl), C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkyl with one double bond and one triple bond, -R₁₁₀, -R₁₂₀, or C₁-C₆ alkyl optionally substituted with 1, 2, 3, or 4 groups independently selected from -OH, -NH₂, C₁-C₃ alkoxy, R₁₁₀, and halogen;
- R_{150'} is C₃-C₇ cycloalkyl, -(C₁-C₃ alkyl)-(C₃-C₇ cycloalkyl), C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ alkyl with one double bond and one triple bond, -R₁₁₀, -R₁₂₀, or C₁-C₆ alkyl optionally substituted with 1, 2, 3, or 4 groups independently selected from -OH, -NH₂, C₁-C₃ alkoxy, R₁₁₀, and halogen;
- R₁₈₀ is selected from morpholinyl, thiomorpholinyl, piperazinyl, piperidinyl, homomorpholinyl, homothiomorpholinyl, homothiomorpholinyl S-oxide, homothiomorpholinyl S,S-dioxide, pyrrolinyl and pyrrolidinyl, each of which is optionally substituted with 1, 2, 3, or 4 groups independently selected from C₁-C₆ alkyl, C₁-C₆ alkoxy, halogen, hydroxy, cyano, nitro, amino, mono(C₁-C₆)alkylamino, di(C₁-C₆)alkylamino, C₂-C₆ alkenyl, C₂-C₆ alkynyl, C₁-C₆ haloalkyl, C₁-C₆ haloalkoxy, amino(C₁-C₆)alkyl, mono(C₁-C₆)alkylamino(C₁-C₆)alkyl, di(C₁-C₆)alkylamino(C₁-C₆)alkyl, and =O;

R_{110} is aryl optionally substituted with 1 or 2 R_{125} groups;
 R_{125} at each occurrence is independently halogen, amino, mono- or dialkylamino, -OH, -C≡N, -SO₂-NH₂, -SO₂-NH-C₁-C₆ alkyl, -SO₂-N(C₁-C₆ alkyl)₂, -SO₂-(C₁-C₄ alkyl), -CO-NH₂, -CO-NH-C₁-C₆ alkyl, or -CO-N(C₁-C₆ alkyl)₂, or
C₁-C₆ alkyl, C₂-C₆ alkenyl or C₂-C₆ alkynyl, each of which is optionally substituted with 1, 2, or 3 groups that are independently selected from C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, amino, and mono- and dialkylamino, or
C₁-C₆ alkoxy optionally substituted with one, two or three of halogen;
 R_{120} is heteroaryl, which is optionally substituted with 1 or 2 R_{125} groups; and
 R_{130} is heterocyclyl optionally substituted with 1 or 2 R_{125} groups; and
 R_2 is ~~selected from the group consisting of H; and C₁-C₆ alkyl, optionally substituted with 1, 2, or 3 substituents that are independently selected from the group consisting of C₁-C₃ alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₃ alkoxy, and -NR₁₋₃, wherein~~
 ~~R_{1-a} and R_{1-b} are H or C₁-C₆ alkyl;~~
~~(CH₂)₃₋₄-aryl; (CH₂)₃₋₄-heteroaryl; C₂-C₆ alkenyl; C₂-C₆ alkynyl; -CONR₁₋₂R₁₋₃; -SO₂NR₁₋₂R₁₋₃; -CO₂H; and -CO₂-(C₁-C₄ alkyl);~~

~~R₃ is selected from the group consisting of H, C₁-C₆-alkyl, optionally substituted with 1, 2, or 3 substituents independently selected from the group consisting of C₁-C₆-alkyl, halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆-alkoxy, and -NR₁₋₂, R₁₋₂ = -(CH₂)₀₋₄-aryl, -(CH₂)₀₋₄-heteroaryl, C₂-C₆-alkenyl, C₂-C₆-alkynyl, -CO-NR₁₋₂-R₁₋₂, -SO₂-NR₁₋₂-R₁₋₂, -CO₂H, and -CO-O-(C₁-C₄-alkyl),~~

~~wherein~~

~~R_{N-2} and R_{N-2} at each occurrence are independently selected from the group consisting of C₁-C₆-alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of -OH, -NH₂, phenyl and halogen, C₆-C₆-cycloalkyl, -(C₁-C₆-alkyl)-(C₆-C₆-cycloalkyl), -(C₁-C₆-alkyl)-O-(C₁-C₆-alkyl), C₂-C₆-alkenyl, C₂-C₆-alkynyl, C₂-C₆-alkyl chain with one double bond and one triple bond, aryl, heteroaryl, heterocycloalkyl, or~~

~~R_{N-2}, R_{N-2} and the nitrogen to which they are attached form a 5, 6, or 7 membered heterocycloalkyl or heteroaryl group, wherein said heterocycloalkyl or heteroaryl group is optionally fused to a benzene, pyridine, or pyrimidine ring, and said groups are unsubstituted or substituted with 1, 2, 3, 4, or 5 groups that at each occurrence are independently C₁-C₆-alkyl, C₁-C₆-alkoxy, halogen, halo C₁-C₆-alkyl, halo C₁-C₆-alkoxy, -CN, -NO₂, -NH₂, NH(C₁-C₆-alkyl), N(C₁-C₆-alkyl)(C₁-C₆-alkyl), -OH, -C(O)NH₂, -C(O)NH(C₁-C₆-alkyl), -C(O)N(C₁-C₆-alkyl)(C₁-C₆-alkyl), C₁-C₆-alkoxy, C₂-C₆-alkyl, C₁-C₆-thioalkoxy, and C₁-C₆-thioalkoxy C₁-C₆-alkyl,~~

~~or wherein~~

~~R₂, R₃ and the carbon to which they are attached form a carbocycle of three thru seven carbon atoms, wherein one carbon~~

~~atom is optionally replaced by a group selected from -O-, -S-,
-SO₂-, or -NR₁₀₄-.~~

2-4. (Canceled)

5. (Currently Amended) A compound according to claim ~~4~~ 1,
wherein R₁ is
-C₁-C₆ alkyl-aryl, -C₁-C₆ alkyl-heteroaryl, or -C₁-C₆ alkyl-
heterocyclyl, where the ring portions of each are optionally
substituted with 1, 2, 3, or 4 groups independently selected from
halogen, -OH, -SH, -C≡N, -NO₂, -NR₁₀₅R'₁₀₅, -CO₂R, -N(R)COR', ~~or~~
~~-N(R)SO₂R' (where R₁₀₅, R'₁₀₅, R and R' are as defined above),~~
-C(=O)-(C₁-C₄) alkyl, -SO₂-amino, -SO₂-mono or dialkylamino,
-C(=O)-amino, -C(=O)-mono or dialkylamino, -SO₂-(C₁-C₄) alkyl, or
C₁-C₆ alkoxy optionally substituted with 1, 2, or 3 groups
which are independently selected from halogen, ~~or~~

~~C₃-C₄ cycloalkyl optionally substituted with 1, 2, or 3
groups independently selected from halogen, -OH, -SH, -C≡N,
-CF₃, C₁-C₆ alkoxy, amino, -C₁-C₆ alkyl and mono- or dialkylamino,
or~~

~~C₁-C₆ alkyl optionally substituted with 1, 2, or 3 groups
independently selected from halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆
alkoxy, amino, mono- or dialkylamino and -C₁-C₂ alkyl, or~~

~~C₂-C₁₀ alkenyl or C₂-C₁₀ alkynyl each of which is optionally
substituted with 1, 2, or 3 groups independently selected from
halogen, -OH, -SH, -C≡N, -CF₃, C₁-C₆ alkoxy, amino, C₁-C₆ alkyl
and mono- or dialkylamino, and the heterocyclyl group is
optionally further substituted with oxo.~~

6. (Original) A compound according to claim 1 wherein:
R_N is -C(=O)-R₁₀₆; and

R₁₀₀ represents aryl, or heteroaryl, where the ring portions of each are optionally substituted with 1, 2, or 3 groups independently selected from

-OR, -NO₂, C₁-C₆ alkyl, halogen, -C≡N, -OCF₃, -CF₃, -(CH₂)₀₋₄-O-P(=O)(OR)(OR'), -(CH₂)₀₋₄-CO-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-O-(CH₂)₀₋₄-CONR₁₀₂R₁₀₂', -(CH₂)₀₋₄-CO-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkenyl), -(CH₂)₀₋₄-CO-(C₂-C₁₂ alkynyl), -(CH₂)₀₋₄-CO-(CH₂)₀₋₆(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-R₁₁₀, -(CH₂)₀₋₄-R₁₂₀, -(CH₂)₀₋₄-R₁₃₀, -(CH₂)₀₋₄-CO-R₁₁₀, -(CH₂)₀₋₄-CO-R₁₂₀, -(CH₂)₀₋₄-CO-R₁₃₀, -(CH₂)₀₋₄-CO-R₁₄₀, -(CH₂)₀₋₄-CO-O-R₁₅₀, -(CH₂)₀₋₄-SO₂-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-SO-(C₁-C₈ alkyl), -(CH₂)₀₋₄-SO₂-(C₁-C₁₂ alkyl), -(CH₂)₀₋₄-SO₂-(CH₂)₀₋₄-(C₃-C₇ cycloalkyl), -(CH₂)₀₋₄-N(R₁₅₀)-CO-O-R₁₅₀, -(CH₂)₀₋₄-N(R₁₅₀)-CO-N(R₁₅₀)₂, -(CH₂)₀₋₄-N(R₁₅₀)-CS-N(R₁₅₀)₂, -(CH₂)₀₋₄-N(R₁₅₀)-CO-R₁₀₅, -(CH₂)₀₋₄-NR₁₀₅R'₁₀₅, -(CH₂)₀₋₄-R₁₄₀, -(CH₂)₀₋₄-O-CO-(C₁-C₆ alkyl), -(CH₂)₀₋₄-O-P(O)-(O-R₁₁₀)₂, -(CH₂)₀₋₄-O-CO-N(R₁₅₀)₂, -(CH₂)₀₋₄-O-CS-N(R₁₅₀)₂, -(CH₂)₀₋₄-O-(R₁₅₀), -(CH₂)₀₋₄-O-R₁₅₀'-COOH, -(CH₂)₀₋₄-S-(R₁₅₀), -(CH₂)₀₋₄-N(R₁₅₀)-SO₂-R₁₀₅, -(CH₂)₀₋₄-C₃-C₇ cycloalkyl, (C₂-C₁₀) alkenyl, or (C₂-C₁₀) alkynyl.

7. (Currently Amended) A compound according to claim 1 wherein:

R_C is hydrogen, -(CR₂₄₅R₂₅₀)₀₋₄-aryl, -(CR₂₄₅R₂₅₀)₀₋₄-heteroaryl, -(CR₂₄₅R₂₅₀)₀₋₄-heterocyclyl, C₆-C₁₄ alkyl optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, R₁₄₀, R₁₂₀, R₁₃₀, -OC=ONR₂₃₅R₂₄₀, -S(=O)₀₋₂(C₁-C₆ alkyl), -SH, and -S(=O)₂NR₂₃₅R₂₄₀, -(CH₂)₀₋₃-(C₆-C₆) cycloalkyl wherein the cycloalkyl is optionally substituted with 1, 2, or 3 groups independently selected from the group consisting of R₂₀₅, -CO₂H, and -CO₂-(C₁-C₄ alkyl), or

~~C₂-C₁₀-alkenyl or C₂-C₁₀-alkynyl, each of which is optionally substituted with 1, 2, or 3 independently selected R₂₀₀ groups, wherein~~

each aryl and heteroaryl is optionally substituted with 1, 2, or 3 R₂₀₀, and wherein each heterocyclyl is optionally substituted with 1, 2, 3, or 4 independently selected R₂₁₀ groups.

8. (Canceled)

9. (Previously Presented) A compound according to claim 1 selected from the group consisting of:

N-(3,5-difluorobenzyl)-N-((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)-5-methyl-N',N'-dipropylisophthalamide;

N-[2-(3,5-difluorophenyl)ethyl]-N-((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)-5-methyl-N',N'-dipropylisophthalamide;

3-[[2-(3,5-difluorophenyl)ethyl]((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)amino)sulfonyl]-N,N-dipropylbenzamide;

N-(3,5-difluorobenzyl)-N-((2R)-3-[[4(R)-6-ethyl-2,2-dioxido-3,4-dihydro-1H-isothiochromen-4-yl]amino]-2-hydroxypropyl)-5-methyl-N',N'-dipropylisophthalamide;

N-[2-(3,5-difluorophenyl)ethyl]-N-((2R)-3-[[4(R)-6-ethyl-2,2-dioxido-3,4-dihydro-1H-isothiochromen-4-yl]amino]-2-hydroxypropyl)-5-methyl-N',N'-dipropylisophthalamide;

3-[[2-(3,5-difluorophenyl)ethyl]((2R)-3-[[4(R)-6-ethyl-2,2-dioxido-3,4-dihydro-1H-isothiochromen-4-yl]amino]-2-hydroxypropyl)amino)sulfonyl]-N,N-dipropylbenzamide;

N-(3,5-difluorobenzyl)-N-((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)-N',N',5-trimethylisophthalamide;

N-[2-(3,5-difluorophenyl)ethyl]-N-((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)-N',N',5-trimethylisophthalamide;

3-[[2-(3,5-difluorophenyl)ethyl]((2R)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)amino)sulfonyl]-N,N-dimethylbenzamide;

N - (3,5-difluorobenzyl)- N - ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)- N' , N' , 5-trimethylisophthalamide;

N -[2-(3,5-difluorophenyl)ethyl]- N - ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)- N' , N' , 5-trimethylisophthalamide;

3-{[[2-(3,5-difluorophenyl)ethyl] ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)amino]sulfonyl}- N , N -dimethylbenzamide;

N - (3-chloro-5-fluorobenzyl)- N -{ (2*R*)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl}-5-methyl- N' , N' -dipropylisophthalamide;

N -[2-(3-chloro-5-fluorophenyl)ethyl]- N -{ (2*R*)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl}-5-methyl- N' , N' -dipropylisophthalamide;

3-[[2-(3-chloro-5-fluorophenyl)ethyl]{ (2*R*)-2-hydroxy-3-[(3-iodobenzyl)amino]propyl)amino)sulfonyl]- N , N -dipropylbenzamide;

N - (3-chloro-5-fluorobenzyl)- N - ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)-5-methyl- N' , N' -dipropylisophthalamide;

N -[2-(3-chloro-5-fluorophenyl)ethyl]- N - ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)-5-methyl- N' , N' -dipropylisophthalamide;

3-{[[2-(3-chloro-5-fluorophenyl)ethyl] ((2*R*)-3-{[(4*R*)-6-ethyl-2,2-dioxido-3,4-dihydro-1*H*-isothiochromen-4-yl]amino}-2-hydroxypropyl)amino]sulfonyl}- N , N -dipropylbenzamide;

N -[(2*R*)-3-(benzylamino)-2-hydroxypropyl]- N - (3,5-difluorobenzyl)-5-methyl- N' , N' -dipropylisophthalamide;

N -[(2*R*)-3-(benzylamino)-2-hydroxypropyl]- N -[2-(3,5-difluorophenyl)ethyl]-5-methyl- N' , N' -dipropylisophthalamide;

3-([[(2*R*)-3-(benzylamino)-2-hydroxypropyl][2-(3,5-difluorophenyl)ethyl]amino)sulfonyl]- N , N -dipropylbenzamide; and salts thereof.

10. (Previously Presented) A pharmaceutical composition comprising a compound according to claim 1, in combination with a physiologically acceptable carrier or excipient.

11-12. (Canceled)

13. (Withdrawn) A method for treating a patient who has, or in preventing a patient from getting, a disease or condition selected from the group consisting of Alzheimer's disease, for helping prevent or delay the onset of Alzheimer's disease, for treating patients with mild cognitive impairment (MCI) and preventing or delaying the onset of Alzheimer's disease in those who would progress from MCI to AD, for treating Down's syndrome, for treating humans who have Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, for treating cerebral amyloid angiopathy and preventing its potential consequences, i.e. single and recurrent lobar hemorrhages, for treating other degenerative dementias, including dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical basal degeneration, or diffuse Lewy body type of Alzheimer's disease and who is in need of such treatment, comprising administering to such patient a therapeutically effective amount of a compound of claim 1.

14. (Withdrawn) A method for the treatment or prevention of Alzheimer's disease, mild cognitive impairment Down's syndrome, Hereditary Cerebral Hemorrhage with Amyloidosis of the Dutch-Type, cerebral amyloid angiopathy, other degenerative dementias, dementias of mixed vascular and degenerative origin, dementia associated with Parkinson's disease, dementia associated with progressive supranuclear palsy, dementia associated with cortical

basal degeneration, diffuse Lewy body type of Alzheimer's disease comprising administration of a therapeutically effective amount of a compound or salt according to Claim 1, to a patient in need thereof.

15. (Withdrawn) A method for making a compound of claim 1.